- 1. A process for the preparation of a controlled release oral dosage form comprising:
- (a) forming granules comprising oxycodone hydrochloride, alkyl cellulose and polymethacrylate, and
  - (b) drying said granules.
- 2. The process of claim 1, further comprising adding aliphatic alcohol and regranulating and compressing said granules into tablets.
- 3. The process of claim 1, wherein said granules are dried at  $50^{\circ}\text{C}$ .
- 4. A process for the preparation of a controlled release oral dosage form comprising:
- (a) forming spheroids comprising oxycodone hydrochloride, spheronising agent, alkyl cellulose and polymethacrylate and

drying said spheroids

- 5. The process of claim 4, where the spheronising agent is microcrystalline cellulose.
- 6. The process of claim 4, further comprising film coating said spheroids.
- 7. A process for the preparation of a controlled release oral dosage form comprising:
- (a) wet granulating oxycodone hydrochloride, alkyl cellulose and polymethacrylate to form granules of said oxycodone hydrochloride,
  - (b) drying said granules,
  - (c) adding aliphatic alcohol,
- (d) regranulating and compressing said granules into tablets.